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| APPLICATION NO. | FILING DATE | FIRST NAMED INVENTOR | ATTORNEY DOCKET NO. | CONFIRMATION NO. |
|---------------------------------|--------------------------------------|----------------------|---------------------|------------------|
| 10/521,140 | 10/31/2005 | Eva Kontsekova | SONN:065US | 5448 |
| | 7590 08/17/2007 & JAWORSKI L.L.P. | | EXAMINER | |
| 600 CONGRESS AVE. SUITE 2400 | | | CHERNYSHEV, OLGA N | |
| AUSTIN, TX 78701 | | | ART UNIT | PAPER NUMBER |
| | | | 1649 | |
| | | | | |
| | | | MAIL DATE | DELIVERY MODE |
| | | | 08/17/2007 | PAPER |

Please find below and/or attached an Office communication concerning this application or proceeding.

The time period for reply, if any, is set in the attached communication.

| | Application No. | Applicant(s) | | | |
|---|---|---|--|--|--|
| | 10/521,140 | KONTSEKOVA, EVA | | | |
| Office Action Summary | Examiner | Art Unit | | | |
| | Olga N. Chernyshev | 1649 | | | |
| The MAILING DATE of this communication ap | pears on the cover sheet with | h the correspondence address | | | |
| A SHORTENED STATUTORY PERIOD FOR REPL WHICHEVER IS LONGER, FROM THE MAILING D - Extensions of time may be available under the provisions of 37 CFR 1. after SIX (6) MONTHS from the mailing date of this communication. - If NO period for reply is specified above, the maximum statutory period - Failure to reply within the set or extended period for reply will, by statut Any reply received by the Office later than three months after the mailin earned patent term adjustment. See 37 CFR 1.704(b). | DATE OF THIS COMMUNIC 136(a). In no event, however, may a re- will apply and will expire SIX (6) MONT e, cause the application to become ABA | ATION. ply be timely filed HS from the mailing date of this communication. NDONED (35 U.S.C. § 133). | | | |
| Status | | | | | |
| 1) Responsive to communication(s) filed on 11 J | <u>une 2007</u> . | | | | |
| 2a) This action is FINAL . 2b) ☑ This | This action is FINAL. 2b) This action is non-final. | | | | |
| 3) Since this application is in condition for allowa | Since this application is in condition for allowance except for formal matters, prosecution as to the merits is | | | | |
| closed in accordance with the practice under | Ex parte Quayle, 1935 C.D. | 11, 453 O.G. 213. | | | |
| Disposition of Claims | · | | | | |
| 4) ⊠ Claim(s) 24-34 is/are pending in the application 4a) Of the above claim(s) 26-28 and 31-34 is/a 5) ☐ Claim(s) is/are allowed. 6) ☒ Claim(s) 24,29 and 30 is/are rejected. 7) ☐ Claim(s) is/are objected to. 8) ☐ Claim(s) are subject to restriction and/or | are withdrawn from consider | ation. | | | |
| Application Papers | | • | | | |
| 9) ☐ The specification is objected to by the Examine 10) ☑ The drawing(s) filed on 12 January 2005 is/are Applicant may not request that any objection to the Replacement drawing sheet(s) including the correct 11) ☐ The oath or declaration is objected to by the E | e: a)⊠ accepted or b)□ ob drawing(s) be held in abeyand ction is required if the drawing(s | e. See 37 CFR 1.85(a). i) is objected to. See 37 CFR 1.121(d). | | | |
| Priority under 35 U.S.C. § 119 | | · | | | |
| 12) Acknowledgment is made of a claim for foreign a) All b) Some * c) None of: 1. Certified copies of the priority document 2. Certified copies of the priority document 3. Copies of the certified copies of the priority application from the International Bureat * See the attached detailed Office action for a list | ts have been received. ts have been received in Ap prity documents have been r uu (PCT Rule 17.2(a)). | plication No eceived in this National Stage | | | |
| Attachment(s) | | | | | |
| Notice of References Cited (PTO-892) Notice of Draftsperson's Patent Drawing Review (PTO-948) Information Disclosure Statement(s) (PTO/SB/08) Paper No(s)/Mail Date 10/31/5. | Paper No(s). 5) Notice of Inf | Immary (PTO-413) /Mail Date ormal Patent Application ence alignment, 2 pages. | | | |

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DETAILED ACTION

Election/Restrictions

1. Applicant's election with traverse of Group I in the reply filed on June 11, 2007 is acknowledged. The traversal is on the ground(s) that the restriction requirement fails to establish that the groups I and II do not relate to a single inventive concept. This is not found persuasive because the restriction requirement with respect to unity of invention and PCT Rule 13.1 was fully explained at pp. 2-3 of Paper mailed on April 11, 2007. Briefly, inventions of Groups I and II encompass different products (a protein and a transgenic animal), and as such they are not among the combination of categories, which support the unity of invention. Moreover, Applicant's own citation of prior art serves as a reference that the fragments of tau were known before the filing of the instant application, see IDS flied on October 31, 2005. Applicant is further advised that claim 33 is an improper dependent claim as it fails to limit the subject matter of the base claim 24, see 37 CFR 1.75(c), and the "Infringement Test" for dependent claims in MPEP § 608.01(n). Therefore, the fact the claim 33 recites the subject matter of claim 24 cannot serve as a basis for rejoining Groups I and II.

Applicant is further advised that truncated tau molecules of four different types have different structure, require separate sequence search and are not obvious over each other; therefore, the election of species for the purpose of examination is proper. Applicant is reminded that if the generic claim is found to be allowable, all species would be examined.

The requirement is still deemed proper and is therefore made FINAL.

2. Claims 26-28 and 31-34 are withdrawn from further consideration pursuant to 37 CFR 1.142(b), as being drawn to a nonelected invention, there being no allowable generic or linking

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claim. Applicant timely traversed the restriction (election) requirement in the reply filed on June 11, 2007.

3. Claims 24 and 29-30, in so far as they are drawn to a type IIA truncated tau molecule, are under examination in the instant office action.

Claim Rejections - 35 USC § 102

4. The following is a quotation of the appropriate paragraphs of 35 U.S.C. 102 that form the basis for the rejections under this section made in this Office action:

A person shall be entitled to a patent unless -

(b) the invention was patented or described in a printed publication in this or a foreign country or in public use or on sale in this country, more than one year prior to the date of application for patent in the United States.

5. Claims 24 and 29-30 are rejected under 35 U.S.C. 102(b) as being anticipated by Ghetti et al., WO9962548, 1999.

Claims 24 and 29-30 encompass an isolated tau molecule comprising the amino acid sequence of SEQ ID NO: 15. Document of Ghetti et al. discloses a form of tau protein, which has an amino acid sequence 100% identical to the instant claimed peptide of SEQ ID NO: 15 (see copy of the sequence alignment attached to the instant office action), thus fully anticipating the instant invention.

Conclusion

6. No claim is allowed.

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Any inquiry concerning this communication or earlier communications from the examiner should be directed to Olga N. Chernyshev whose telephone number is (571) 272-0870. The examiner can normally be reached on 8:00 AM to 5:00 PM.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Christina Y. Chan can be reached on (571) 272-0841. The fax phone number for the organization where this application or proceeding is assigned is 571-273-8300.

Information regarding the status of an application may be obtained from the Patent Application Information Retrieval (PAIR) system. Status information for published applications may be obtained from either Private PAIR or Public PAIR. Status information for unpublished applications is available through Private PAIR only. For more information about the PAIR system, see http://pair-direct.uspto.gov. Should you have questions on access to the Private PAIR system, contact the Electronic Business Center (EBC) at 866-217-9197 (toll-free). If you would like assistance from a USPTO Customer Service Representative or access to the automated information system, call 800-786-9199 (IN USA OR CANADA) or 571-272-1000.

Olga N. Chernyshev, Ph.D.

Primary Examiner Art Unit 1649

August 14, 2007

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<!--StartFragment-->RESULT 7
AAY15200
     AAY15200 standard; protein; 351 AA.
XX
     AAY15200;
AC
XX
DT
     28-FEB-2000
                  (first entry)
XX
DE
     Human Tau protein.
XX
KW
     Human Tau gene; neurofibrillary tangle formation; abnormal tau filament;
     brain; mutation; phosphorylation; isoform ratio; diagnosis; tauopathy;
KW
KW
     treatment; neurodegenerative disorder; Fronto-Temporal Dementia;
KW
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KW
     Pick's Disease; Progressive Supranuclear Palsy; PSP; Alzheimer's Disease;
KW
     Corticobasal Degeneration; CD; Prion Protein Cerebral Amyloid Angiopathy;
KW
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XX
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XX
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FT
                     /note= "Microtubule related protein"
XX
PN
     W09962548-A1.
XX
·PD
     09-DEC-1999.
XX
PF
     28-MAY-1999;
                    99WO-US012036.
XX
PR
     01-JUN-1998:
                    98US-0087557P.
XX
PA
     (ADRE-) ADVANCED RES & TECHNOLOGY INST.
\mathbf{x}\mathbf{x}
PI
     Ghetti B,
               Spillantini MG, Murrell JR,
                                             Goedert M.
PI
     Klug A;
XX
DR
     WPI; 2000-086858/07.
DR
     N-PSDB; AAZ29262.
XX
     Diagnosing a tauopathy, especially a Fronto-Temperal Dementia.
PT
XX
PS
     Disclosure; Page 85; 90pp; English.
XX
CC
     The present amino acid sequence is a form of human Tau protein. There are
CC
     six tau isoforms, expressed in the normal adult brain with a slight
     preponderance of those with 3 repeats over those with 4 repeats.
CC
     Mutations in the tau gene affects phosphorylation and leads to formation
     of neurofibrillary tangles and alters the tau isoform ratio. The
CC
CC
     increased ratio of 4:3 repeat and abnormal tau filaments is closely
CC
     related to neurodegenerative disorders. This sequence can be used for
     diagnosis and treatment of tauopathies, like Fronto-Temporal Dementia,
CC
CC
     Familial Multiple System Tauopathy with presentle Dementia (MSTD), Pick's
CC
     Disease, Progressive Supranuclear Palsy (PSP), Corticobasal Degeneration
     (CD) or Alzheimer's Disease. A composition that decreases the ratio of
CC
     4:3 repeat tau isomers, along with an agent for treatment of a cognitive
CC
     disorder, is useful for treating a tauopathy. It may also be useful in
CC
     diagnosis of Prion Protein Cerebral Amyloid Angiopathy and other prion
CC
     protein associated disease characterized by abnormal tau filament
CC
     formation
xx
SO
     Sequence 351 AA;
  Query Match
                         رز 100.08
                                  Score 1108; DB 3; Length 351;
  Best Local Similarity
                          100.0%; Pred. No. 3.8e-76;
  Matches 210; Conservative
                                0; Mismatches
                                                      Indels
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Qÿ
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           93 IATPRGAAPPGQKGQANATRIPAKTPPAPKTPPSSGEPPKSGDRSGYSSPGSPGTPGSRS 152
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